Listing of the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

Claims 1-2 (Canceled)

Claim 3 (**Previously Presented**) The method of Claim 9 or 10, wherein the host cell is a prokaryotic cell or an eukaryotic cell.

Claim 4 (**Previously Presented**) The method of Claim 3, wherein the host cell is a microorganism.

Claim 5 (**Previously Presented**) The method of Claim 4, wherein the microorganism is *Escherichia coli*.

Claim 6 (**Currently Amended**) The method of Claim 9 or Claim 10, wherein the molecular weight of the polypeptide comprising a serine residue is about 1000 to 20000 <u>daltons</u>.

Claim 7 (Canceled)

Claim 8 (**Previously Presented**) The method of Claim 9 or Claim 10, wherein the atrial natriuretic peptide is human atrial natriuretic peptide.

Claim 9 (**Currently Amended**) A method for reducing formation of a byproduct polypeptide comprising an O-acetylserine residue in place of a serine residue in the atrial natriuretic peptide, comprising:

- (i) culturing, in a medium, transformed host cells that produce a recombinant atrial natriuretic peptide comprising a serine residue and a byproduct polypeptide comprising an O-acetylserine residue in place of a serine residue;
- (ii) adding <u>at least 3 g/L</u> methionine and at least one of <u>at least 3 g/L</u> histidine or <u>at least 3 g/L</u> glycine to the medium in an amount effective to reduce said byproduct formation; and
- (iii) reducing the formation of said byproduct polypeptide,
 wherein the formation of said byproduct polypeptide is reduced in an
 amount greater than or equal to 50% as compared to a control medium
 with no methionine, histidine, or glycine added.

Claim 10 (**Currently Amended**) A method for producing a polypeptide comprising a serine residue comprising:

- (i) culturing, in a medium, transformed host cells that produce a recombinant atrial natriuretic peptide comprising a serine residue and a byproduct polypeptide comprising an O-acetylserine residue in place of a serine residue in the atrial natriuretic peptide;
- (ii) adding <u>at least 3 g/L</u> methionine and at least one of <u>at least 3 g/L</u> histidine or <u>at least 3 g/L</u> glycine to the medium in an amount effective to reduce said byproduct formation; and
- (iii) reducing the formation of said byproduct polypeptide,
 wherein the formation of said byproduct polypeptide is reduced in an
 amount greater than or equal to 50% as compared to a control medium
 with no methionine, histidine, or glycine added.

Claims 11-16 (Canceled)

Claim 17 (**Currently Amended**) A method for reducing formation of a byproduct polypeptide comprising an O-acetylserine residue in place of a serine residue <u>in the atrial natriuretic peptide</u>, comprising:

- (i) culturing, in a medium, transformed host cells that produce a recombinant atrial natriuretic peptide comprising a serine residue and a byproduct polypeptide comprising an O-acetylserine residue in place of a serine residue;
- (ii) adding to said medium at least one of <u>at least 3 g/L</u> histidine or <u>at least 3 g/L</u> glycine to the medium in an amount effective to reduce said byproduct formation; and
- (iii) reducing the formation of said byproduct polypeptide,
 wherein the formation of said byproduct polypeptide is reduced in an
 amount greater than or equal to 50% as compared to a control medium
 with no histidine or glycine added.

Claim 18 (**Currently Amended**) A method for producing a polypeptide comprising a serine residue comprising:

- (i) culturing, in a medium, transformed host cells that produce a recombinant atrial natriuretic peptide comprising a serine residue and a byproduct polypeptide comprising an O-acetylserine residue in place of a serine residue in the atrial natriuretic peptide;
- (ii) adding at least one of <u>at least 3 g/L</u> histidine or <u>at least 3 g/L</u> glycine to the medium in an amount effective to reduce said byproduct formation; and
- (iii) reducing the formation of said byproduct polypeptide,
 wherein the formation of said byproduct polypeptide is reduced in an
 amount greater than or equal to 50% as compared to a control medium
 with no histidine or glycine added.

Claims 19-20 (Canceled)

Claim 21 (**Previously Presented**) The method of Claim 17, wherein the host cell is a prokaryotic cell or an eukaryotic cell.

Claim 22 (**Previously Presented**) The method of Claim 21, wherein the host cell is a microorganism.

Claim 23 (**Previously Presented**) The method of Claim 22, wherein the microorganism is *Escherichia coli*.

Claim 24 (**Currently Amended**) The method of Claim 17, wherein the molecular weight of the polypeptide comprising a serine residue is about 1000 to 20000 <u>daltons</u>.

Claim 25 (**Previously Presented**) The method of Claim 17, wherein the atrial natriuretic peptide is human atrial natriuretic peptide.

Claim 26 (**Previously Presented**) The method of Claim 17, further comprising adding an the amount of methionine effective to reduce formation of a byproduct polypeptide wherein said amount is 3 g/L.

Claim 27 (**Previously Presented**) The method of Claim 18, wherein the host cell is a prokaryotic cell or an eukaryotic cell.

Claim 28 (**Previously Presented**) The method of Claim 27, wherein the host cell is a microorganism.

Claim 29 (**Previously Presented**) The method of Claim 28, wherein the microorganism is *Escherichia coli*.

Claim 30 (**Currently Amended**) The method of Claim 18, wherein the molecular weight of the polypeptide comprising a serine residue is about 1000 to 20000 <u>daltons</u>.

Claim 31 (**Previously Presented**) The method of Claim 18, wherein the atrial natriuretic peptide is human atrial natriuretic peptide.

PATENT APPLICATION NO. 10/030,452 ATTORNEY DOCKET NO. 58777.000008

Claim 32 (**Previously Presented**) The method of Claim 18, further comprising adding an the amount of methionine effective to reduce formation of a byproduct polypeptide wherein said amount is 3 g/L.

Claims 33-36 (Canceled)